

|                |  |                                 |
|----------------|--|---------------------------------|
| Run on:        | October 1, 1999, 15:34:36<br>(without alignments)  | Search time 16:19.02 Seconds    |
| Scoring table: | IDENTITY_NUC   | 45.180 Million cell updates/sec |
| Searched:      | 679419 seqs,   | 1590154680 residues             |
| Database :     | GenEmbl:<br>1: gb_ba1:<br>2: gb_ba2:<br>3: gb_om:<br>4: gb_ov:<br>5: gb_bp:<br>6: gb_ph:<br>7: gb_dl1:<br>8: gb_dl2:<br>9: gb_pr1:<br>10: gb_pr2:<br>11: gb_pr3:<br>12: gb_ro:<br>13: gb_st:<br>14: gb_sts:<br>15: gb_sy:<br>16: gb_un:<br>17: gb_vl:<br>18: em_fun:<br>19: em_htg:<br>20: em_hum1:<br>21: em_hum2:<br>22: em_in1:<br>23: em_om:<br>24: em_or:<br>25: em_cv:<br>26: em_ph:<br>27: em_p1:<br>28: em_p2:<br>29: em_xo:<br>30: em_sts:<br>31: em_sy:<br>32: em_un:<br>33: em_vl:<br>34: gb_htg1:<br>35: gb_htg2:<br>36: gb_in1:<br>37: gb_in2:<br>38: em_ba1:<br>39: em_ba2:<br>40: em_hum3:<br>41: em_hum4:<br>42: gb_pr4:<br>43: em_cv2:<br>44: em_cv3:<br>45: em_cv4:<br>46: em_cv5:<br>47: em_cv6:<br>48: em_cv7:<br>49: em_cv8:<br>50: em_cv9:<br>51: em_cv10:<br>52: em_cv11:<br>53: em_cv12:<br>54: em_cv13:<br>55: em_cv14:<br>56: em_cv15:<br>57: em_cv16:<br>58: em_cv17:<br>59: em_cv18:<br>60: em_cv19:<br>61: em_cv20:<br>62: em_cv21:<br>63: em_cv22:<br>64: em_cv23:<br>65: em_cv24:<br>66: em_cv25:<br>67: em_cv26:<br>68: em_cv27:<br>69: em_cv28:<br>70: em_cv29:<br>71: em_cv30:<br>72: em_cv31:<br>73: em_cv32:<br>74: em_cv33:<br>75: em_cv34:<br>76: em_cv35:<br>77: em_cv36:<br>78: em_cv37:<br>79: em_cv38:<br>80: em_cv39:<br>81: em_cv40:<br>82: em_cv41:<br>83: em_cv42:<br>84: em_cv43:<br>85: em_cv44:<br>86: em_cv45:<br>87: em_cv46:<br>88: em_cv47:<br>89: em_cv48:<br>90: em_cv49:<br>91: em_cv50:<br>92: em_cv51:<br>93: em_cv52:<br>94: em_cv53:<br>95: em_cv54:<br>96: em_cv55:<br>97: em_cv56:<br>98: em_cv57:<br>99: em_cv58:<br>100: em_cv59:<br>101: em_cv60:<br>102: em_cv61:<br>103: em_cv62:<br>104: em_cv63:<br>105: em_cv64:<br>106: em_cv65:<br>107: em_cv66:<br>108: em_cv67:<br>109: em_cv68:<br>110: em_cv69:<br>111: em_cv70:<br>112: em_cv71:<br>113: em_cv72:<br>114: em_cv73:<br>115: em_cv74:<br>116: em_cv75:<br>117: em_cv76:<br>118: em_cv77:<br>119: em_cv78:<br>120: em_cv79:<br>121: em_cv80:<br>122: em_cv81:<br>123: em_cv82:<br>124: em_cv83:<br>125: 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Clones are available from the UK MRC Human Genome Mapping Project Resource Centre, Hinckley, Leicestershire CB10 1RQ, UK. See URL: <http://www.hgmp.mrc.ac.uk/> for details or contact: biocenter@hgmp.mrc.ac.uk.

Location/Qualifiers

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/db\_xref="taxon:9606"  
/sex="male"

/dev\_stage="adult"

/tissue\_type="blood"

/clone\_lib="CGI-1"

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Best Local Similarity 63.6%; Pred. No. 2.8e+02;  
Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

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/transcript="MADPRAIDLPGRARNEPDVYETSLPLDQEDDAFQEELELT

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CDS

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BASE COUNT 473 a 445 c 453 g 350 t  
ORIGIN

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| RESULT 3  |  |   |   |
| LOCUS CEF18E2/c   |  | DEFINITION Caenorhabditis elegans cosmid F18E2, complete sequence.  |   |
| DEFINITION Caenorhabditis elegans   |  | ACCESION 275537   |   |
| ACCESION 275537   |  | NID 91418504  |   |
| NID 91418504  |  | VERSION 275537.1  |   |
| VERSION 275537.1  |  | KEYWORDS GI:1418504<br>HGT; ABC transporter; Acid phosphatase like.   |   |
| KEYWORDS GI:1418504<br>HGT; ABC transporter; Acid phosphatase like.   |  | SOURCE Caenorhabditis elegans   |   |
| SOURCE Caenorhabditis elegans   |  | ORGANISM Caenorhabditis elegans   |   |
| ORGANISM Caenorhabditis elegans   |  | REFERENCE 1 (bases 1 to 25321)  |   |
| REFERENCE 1 (bases 1 to 25321)  |  | AUTHORS Lightning, J.<br>TITLE Direct Submission  |   |
| AUTHORS Lightning, J.<br>TITLE Direct Submission  |  | JOURNAL Submitter (29-JUN-1996) Louis, MO 63110, USA. E-mail: jes@sanjour.ac.uk or rwannematec.wustl.edu  |   |
| JOURNAL Submitter (29-JUN-1996) Louis, MO 63110, USA. E-mail: jes@sanjour.ac.uk or rwannematec.wustl.edu  |  | REFERENCE 2 (bases 1 to 25321)  |   |
| REFERENCE 2 (bases 1 to 25321)  |  | AUTHORS Wilson, R., Ainscough, R., Anderson, K., Baynes, C., Berks, M., Bonfield, J., Burton, J., Cornell, M., Copsey, T., Cooper, J., Coulson, A., Craxton, M., Dear, S., Du, Z., Durbin, R., Favell, A., Fulton, L., Gardner, A., Green, P., Hawkins, J., Laister, N., Hawkins, T., Hillier, L., Johnston, L., Jones, M., Kersting, J., Kirstein, J., Latreille, P., Lightning, J., Lloyd, C., McMurray, A., Mortimor, B., O'Callaghan, M., Parsons, J., Percy, C., Rifkin, L., Saunders, D., Shawken, R., Smalton, N., Smith, A., Sonnhammer, E., Staden, R., Sulston, J., Thierry-Mieg, J., Thomas, K., Vaughn, K., Wilkinson, R., Watson, A., Weinstock, J., Wilkison, R., Watson, J., and Woldman, P. |   |
| AUTHORS Wilson, R., Ainscough, R., Anderson, K., Baynes, C., Berks, M., Bonfield, J., Burton, J., Cornell, M., Copsey, T., Cooper, J., Coulson, A., Craxton, M., Dear, S., Du, Z., Durbin, R., Favell, A., Fulton, L., Gardner, A., Green, P., Hawkins, J., Laister, N., Hawkins, T., Hillier, L., Johnston, L., Jones, M., Kersting, J., Kirstein, J., Latreille, P., Lightning, J., Lloyd, C., McMurray, A., Mortimor, B., O'Callaghan, M., Parsons, J., Percy, C., Rifkin, L., Saunders, D., Shawken, R., Smalton, N., Smith, A., Sonnhammer, E., Staden, R., Sulston, J., Thierry-Mieg, J., Thomas, K., Vaughn, K., Wilkinson, R., Watson, A., Weinstock, J., Wilkison, R., Watson, J., and Woldman, P. |  | JOURNAL Nature 368 (6466), 32-38 (1994)   |   |
| JOURNAL Nature 368 (6466), 32-38 (1994)   |  | COMMENT Coding sequences below are predicted from computer analysis, using predictions from Genefinder (P. Green, U. Washington), and other available information.  |   |
| COMMENT Coding sequences below are predicted from computer analysis, using predictions from Genefinder (P. Green, U. Washington), and other available information.  |  | JOURNAL Nature 368 (6466), 32-38 (1994)   |   |
| JOURNAL Nature 368 (6466), 32-38 (1994)   |  | SEE-SEE: http://webspace.sanger.ac.uk/cgi-bin/display?db=wormbase&class=Sequence &object=F18E2  |   |
| SEE-SEE: http://webspace.sanger.ac.uk/cgi-bin/display?db=wormbase&class=Sequence &object=F18E2  |  | Current sequence finishing criteria for the C. elegans genome sequencing consortium are that all bases are either sequenced unambiguously on both strands, or on a single strand with both a dye primer and dye terminator reaction, from distinct subclones.   |   |
| Current sequence finishing criteria for the C. elegans genome sequencing consortium are that all bases are either sequenced unambiguously on both strands, or on a single strand with both a dye primer and dye terminator reaction, from distinct subclones.   |  | EXCEPTIONS: Exceptions are indicated by an explicit note.   |   |
| EXCEPTIONS: Exceptions are indicated by an explicit note.   |  | IMPORTANT: This sequence is NOT necessarily the entire insert of the specified clone. It may be shorter because we only sequence overlapping sections once, or longer because we arrange for a small overlap between neighbouring submissions.  |   |
| IMPORTANT: This sequence is NOT necessarily the entire insert of the specified clone. It may be shorter because we only sequence overlapping sections once, or longer because we arrange for a small overlap between neighbouring submissions.  |  | IMPORTANT: This sequence is not the entire insert of clone F18E2. It may be shorter because we only sequence overlapping sections once, or longer because we arrange for a small overlap between neighbouring submissions.  |   |
| IMPORTANT: This sequence is not the entire insert of clone F18E2. It may be shorter because we only sequence overlapping sections once, or longer because we arrange for a small overlap between neighbouring submissions.  |  | The true left end of clone F18E2 is at 14866 in the true right end of clone F18E2 is at 14866 in sequence 275554.   |   |
| The true left end of clone F18E2 is at 14866 in the true right end of clone F18E2 is at 14866 in sequence 275554.   |  | The true left end of clone F18E2 is at 25221 in this sequence. The true right end of clone F18E2 is at 24380 in this sequence. The true left end of clone F18E2 is at 25221 in this sequence. The true right end of clone F18E2 is at 24380 in this sequence.   |   |
| The true left end of clone F18E2 is at 25221 in this sequence. The true right end of clone F18E2 is at 24380 in this sequence.  |  | start of this sequence (1..104) overlaps with the end of sequence 275527.   |   |
| start of this sequence (1..104) overlaps with the end of sequence 275527.   |  | The end of this sequence (25221..25321) overlaps with the start of sequence 275554.   |   |

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| /note="Similarity to Aspergillus acid phosphatase (TR:G75244)"  |                                    |  |
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| Best Local Similarity 63.6%; Pred. No. 2.2e+02; Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;   |                                    |  |
| Qy 1 gengtngartggaaattttgcnn 22   |                                    |  |

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| ORGANISM                 | Saccharomyces cerevisiae<br>Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;   |
| REFERENCE                | 1 (bases 1 to 2716)   |
| AUTHORS                  | Gamonet, F. and Lauquin, J.-M.  |
| JOURNAL                  | Unpublished   |
| FEATURES                 | Location/Qualifiers   |
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| REFERENCE                | 2 (bases 1 to 2716)   |
| AUTHORS                  | Gamonet, F. and Lauquin, J.-M.  |
| JOURNAL                  | Unpublished   |
| FEATURES                 | Location/Qualifiers   |
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| RESULT                   | 5   |
| BASE COUNT               | 814 a   |
| ORIGIN                   | 814 a   |
| Query Match              | 67.8%; Score 15.6; DB 3; Length 2061;   |
| Best Local Similarity    | 65.2%; Pred. No. 3.5e+02;   |
| Matches 15; Conservative | 3; Mismatches 5; Indels 0; Gaps 0;  |
| Query                    | 1 gcnqntngartggaaatyggcng 23  |
| Db                       | 1192 GACGTGGAACTGGAAATTCGCTCG 1214  |
| BASE COUNT               | 558 c   |
| ORIGIN                   | 587 g   |
| Query Match              | 67.8%; Score 15.6; DB 7; Length 2716;   |
| Best Local Similarity    | 65.2%; Pred. No. 3.4e+02;   |
| Matches 15; Conservative | 3; Mismatches 5; Indels 0; Gaps 0;  |
| Query                    | 1 gcnqntngartggaaatyggcng 23  |
| Db                       | 1704 GCCGTGAATGAAATTTCGCTCG 1682  |
| RESULT                   | 6   |
| DEFINITION               | SC8419  |
| ACCESSION                | X93502  |
| NID                      | 9108839   |
| VERSION                  | X93502.1  |
| KEYWORDS                 | GI:1089839<br>homociontin<br>bacteri  |
| CONC                     | +   |
| ORGANISM                 | S.cerevisiae lys4 gene.   |
| REFERENCE                | SC8419/C  |
| AUTHORS                  | SC8419  |
| JOURNAL                  | LOCUS   |
| FEATURES                 | DEFINITION  |
| source                   | SC8419  |
| 1. .2716                 | DEFINITION  |
| Organism                 | SC8419  |
| Accession                | SC8419  |
| Version                  | SC8419  |
| Keywords                 | SC8419  |
| Comments                 | SC8419  |
| Origin                   | SC8419  |
| Length                   | 30507 bp  |
| Organism                 | S.cerevisiae chromosome IV cosmid 8419.   |
| Accession                | Z49101  |
| Date                     | 27/12/56  |
| PLN                      | 11-AUG-1997   |







| AUTHORS               | Montell,C.  |
|-----------------------|---|
| TITLE                 | Direct Submission   |
| JOURNAL               | Submitted (12-JUN-1997) C. Montell, The Johns Hopkins University School of Medicine, Department of Biological Chemistry, 725 N. Wolfe Street, Baltimore, MD 21205, USA  |
| COMMENT               | Related sequences: X89068, U47050.  |
| FEATURES              | Location/qualifiers   |
| source                | 1..3448<br>/organism="Homo sapiens"<br>/db_xref="taxon: 9606  |
| gene                  | 425..2971<br>/gene="TRPC3"<br>/codon_start=1  |
| CDS                   | /product="transient receptor potential related channel 3 protein"<br>/protein_id="CAA74083.1"<br>/db_xref="PID:e23535"<br>/db_xref="PID:9225097"<br>/db_xref="GI:2255937"<br>/db_xref="SPTRNBL:Q13507"<br>/translation="MEGSPSLRRMTMREKGRRAVGPAPMENDRGTSLTAAEERFL<br>DAEYGNIPVYRKLMEESKTLVNCYDGMGNALQAVGNHLETELLIKENLARI<br>GDALLIAISKQYVRIYEALNHPGFAASKRUTLSPCEQELDDAYDEGTGTRSPD<br>ITPIILAHCKYEVHMLIKAKGARFPHYECKGDCMKMOKRDSESHRSRINAY<br>KGASPAVLSSLSSEDPLTAELSNEAKLANIEKEFDNRKLSMCQDDEVGVLDL<br>CSDSEVEAINGDLSEAPELVEHRKASLYKATYKTFYAHPNQCOQLITIW<br>YENLSCREOTIAIKCLVLYALGIPFLAGIYWTAPCSRIGKILRSHPNKFVABRAS<br>FILFLGLLIVFNASDRFGCTTLPNTIVDKQIFRNKTCTFTEMILINWVGMW<br>SECKELWLPPETOFTYARDKWNPSDPMOLISELYLAUVLSFSRILQALQATAQQVDSYQES<br>DISEVETULPPEYDFTYARDKWNPSDPMOLISELYLAUVLSFSRILQALQATAQQVDSYQES<br>QISLGRIVKIDFKEFVLMFMPAMIGMELIYSYLGAKUNAAFTVESEPKTFLWS<br>TIGLSEVTISVIVKYDHFETIENGIVLGYIIVVVVLLNLLAMINSSTQEIDDSD<br>VKEFKARSKLWISYFDGKTPPPSFULPSKPSFVYFIMRVYNPFICRRLQDDEM<br>GNGNSKRNLLETSNSRVEFHSENSELNQPTRYQIMKMLIKRVLKAGQYDKENDE<br>VNGELBEIKODISSLRLYELLDRKS9T |
| BASE COUNT            | 945 a<br>772 c<br>802 g<br>929 t  |
| ORIGIN                |   |
| Query Match           | 67.8%   |
| Best Local Similarity | 65.2%   |
| Matches               | 15;   |
| Conservative          | 3;  |
| Mismatches            | 5;  |
| Indels                | 0;  |
| Gaps                  | 0;  |
| RESULT                | 11  |
| AC005154              | AC005154 151630 bp DNA PRI 23-SEP-1998  |
| LOCUS                 | Homo sapiens PAC clone DJ0777023 from 7p14-p15, complete sequence.  |
| DEFINITION            |   |
| AC005154              |   |
| NID                   | 9242763   |
| VERSTON               | AC005154..1 GI:3242763  |
| KEYWORDS              | HTG.  |
| SOURCE                | Human.  |
| ORGANISM              | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  |
| REFERENCE             | Bauer,C., Langston,Y. and Harrison,M.   |
| AUTHORS               | The sequence of Homo sapiens PAC clone Unpublished (1998)   |
| TITLE                 |   |
| JOURNAL               |   |
| AUTHORS               | Waterson,N.   |
| TITLE                 | Direct Submission   |
| JOURNAL               | Submitted (20-JUN-1998) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA  |
| COMMENT               | Submitted By: WUGSC Center Sequencing Center  |

Department of Genetics  
Washington University  
St. Louis MO 63108, USA  
<http://genome.wustl.edu/gsc>  
mailto:[sapiens@watson.wustl.edu](mailto:sapiens@watson.wustl.edu)

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded or sequenced with an alternate chemistry; an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequencing from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:  
The sequence of this clone was established as part of a mapping and sequencing collaboration between the NHGRI Chromosome 7 Mapping Project (Eric D. Green, Director), John D. McPherson in the Department of Genetics (Washington University), and the Washington University Genome Sequencing Center. For additional information about the map position of this sequence, see <http://www.ncbi.nlm.nih.gov/DIR/GTB/CHR7>, send mailto:[egreen@hgri.nih.gov](mailto:egreen@hgri.nih.gov), or see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:  
This clone was derived from human PAC library RPCL-4, prepared by Pieter de Jong and coworkers at Roswell Park Cancer Institute, using the method described by Ioannou et al., Nature Genetics 6:84-9 (1994). The library is from one male donor. For further details, see <http://bacpac.med.buffalo.edu/>. The clone is available from Genome Systems, Inc. (<http://www.genomesystems.com>).

VECTOR: PCYPAC2

NEIGHBORING SEQUENCE INFORMATION:  
The actual start of this clone is at base position 1 of DJ0777023; the actual end is at 151630 of DJ0777023.

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 repeat\_region 7512..7628 /rpt\_family="MIR"  
 STS 8566..8530 /db\_xref="GI:1343978"  
 repeat\_region 9111..9287 /rpt\_family="Alu"  
 repeat\_region 10298..10600 /rpt\_family="Alu"  
 repeat\_region 12876..12895 /rpt\_family="L2"  
 repeat\_region 15146..15465 /rpt\_family="MIR"  
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 repeat\_region 19648..19739 /rpt\_family="MER1\_type"  
 repeat\_region 20111..20419 /rpt\_family="MIR"  
 repeat\_region 20804..20970 /rpt\_family="MER1\_type"  
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 repeat\_region 22552..22979 /rpt\_family="Retroviral"  
 repeat\_region 23039..25680 /rpt\_family="Retroviral"  
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 repeat\_region 27520..27882 /rpt\_family="Retroviral"  
 repeat\_region 27910..28343 /rpt\_family="Retroviral"  
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 Db 86858 GATGTGAGGAAATTGCAAG 86880  
 RESULT 12  
 AF093117 LOCUS AF093117 147216 bp DNA PRI 02-OCT-1998  
 DEFINITION Homo sapiens chromosome 7qtel0 BAC E3, complete sequence.  
 AUTHOR Blechschmidt, K., Nordstiek, G., Drescher, B., Weber, J., Schattevoy, R.,  
 Koerber, J. and Rosenthal, A.  
 TITLE Direct Submission  
 JOURNAL Submitted (18-SEP-1998) Genome Analysis, Institute for Molecular  
 Biotechnology, Beutenbergstrasse 11, Jena 07745, Germany  
 FEATURES Location,Qualifiers  
 SOURCE 1..147216  
 ORGANISM Homo sapiens  
 /organism="Homo sapiens"  
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 REFERENCE 1 (bases 1 to 147216)  
 Blechschmidt, K., Nordstiek, G., Drescher, B., Weber, J., Schattevoy, R.,  
 Koerber, J. and Rosenthal, A.  
 ACCESSION AF093117  
 NID 93657910  
 VERSION AF093117.1 GI:3687910  
 KEYWORDS HTG,  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryote; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 147216)  
 Blechschmidt, K., Nordstiek, G., Drescher, B., Weber, J., Schattevoy, R.,  
 Koerber, J. and Rosenthal, A.  
 TITLE Direct Submission  
 JOURNAL Submitted (18-SEP-1998) Genome Analysis, Institute for Molecular  
 Biotechnology, Beutenbergstrasse 11, Jena 07745, Germany  
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Best Local Similarity 65.2%; Pred. No. 2.3e+02;
Matches 15; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

Qy 1 ggcgttgaaatggaaatggcg 23
Db 107284 GCAGTTGGTGGAACTTGGCAGG 107306

RESULT 13
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LOCUS 3417 bp mRNA channel (http://mRNA_complete.cds.
DEFINITION Human putative calcium influx channel (http://mRNA_complete.cds.
ACCESSION U47050
VERSION 92295902
NID U47050.1 GI:2295902
KEYWORDS human.
SOURCE Homo sapiens
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
REFERENCE 1 (bases 1 to 3417)
AUTHORS Zhu,X., Jiang,M., Peyton,M., Bouley,G., Hurst,R., Stefani,E. and
TITLE Birnbaumer,L.
JOURNAL trp, a novel mammalian gene family essential for agonist-activated
MEDLINE capacitative Ca2+ entry
JOURNAL Cell 85 (5), 661-671 (1996)
REFERENCE 96234226
AUTHORS 2 (bases 1 to 3417)
TITLE Direct Submission
JOURNAL Submitted (24-JUN-1996) Xi Zhu, Anesthesiology, UCLA School of
Medicine, BH-612, CHS, Los Angeles, CA 90095-1778, USA
COMMENT On Aug 4, 1997 this sequence version replaced gi:1336112.
FEATURES Location/Qualifiers
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ORGANISM "Homo sapiens"
AUTHORS /db_xref="Taxon:9606"
TITLE /gene="Htrp3"
JOURNAL /note="Htrp3; putative calcium influx channel; stimulated by agonist to some G protein coupled receptors; similar to Drosophila transient receptor potential" /codon_start_1
COMMENT CDS

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| LOCUS      | AF111107  | 3691 bp  | mRNA       | ROD | 15-MAR-1999 |  |  |  |  |
| DEFINITION | Mus musculus  | transient receptor potential 2 (TRP2) mRNA, complete cds.                    |            |     |             |  |  |  |  |
| ACCESSION  | AF111107  |  |            |     |             |  |  |  |  |
| VERSION    | 94324537  |  |            |     |             |  |  |  |  |
| KEYWORDS   | NID   | AF111107.1   | GI:4324937 |     |             |  |  |  |  |
| SOURCE     | house mouse   |  |            |     |             |  |  |  |  |
| ORGANISM   | Mus musculus  |  |            |     |             |  |  |  |  |
| REFERENCE  | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Rodentia; Sciurognathii; Muridae; Murinae; Mus.             |  |            |     |             |  |  |  |  |
| AUTHORS    | 1 (bases 1 to 3691)   |  |            |     |             |  |  |  |  |
| TITLE      | Vannier,B., Peyron,M., Boulay,G., Brown,D., Qin,N., Jiang,M., Xu,X. and Birnbaumer,L.   |  |            |     |             |  |  |  |  |
| JOURNAL    | Mouse trp2, the homologue of the human trpc2 pseudogene, encodes mTrp2, a store depletion-activated capacitative Ca2+ entry channel |  |            |     |             |  |  |  |  |
| MEDLINE    | Proc. Natl. Acad. Sci. U.S.A. 96 (5), 2060-2064 (1999)  |  |            |     |             |  |  |  |  |
| REFERENCE  | 99162557  |  |            |     |             |  |  |  |  |
| AUTHORS    | 2 (bases 1 to 3691)   |  |            |     |             |  |  |  |  |
| TITLE      | Vannier,B., Peyron,M., Zhu,X. and Birnbaumer,L.   |  |            |     |             |  |  |  |  |
| JOURNAL    | Direct Submission   |  |            |     |             |  |  |  |  |
| JOURNAL    | Submitted: 04-DEC-1998  | Anesthesiology, UCLA, 10833 Le Conte Avenue, Los Angeles, CA 90095-1778, USA |            |     |             |  |  |  |  |

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RES source
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 BASE COUNT 769 a 1091 c 1014 g 817 t  
 ORIGIN







(SUMO ) SUMITOMO CHEM CO LTD.  
PI Mori S, Nakanishi H, Takahashi M;  
WPI: 98-439341/38.  
DR  
PI New nicotianamine aminotransferase protein and DNA - useful for  
enhancing iron absorption of plant cells  
PS Claim 16: Page 7; 17PP; English.  
CC The primers V48149 and V48150 were used in the cloning of nicotianamine  
CC aminotransferase can be used in a plasmid to transform plant cells to  
CC produce cells with enhanced iron absorption, and it is implied [though  
CC not stated] that plants with improved resistance to iron deficiency  
CC chlorosis in calcareous soils can be regenerated from the transformed  
CC cells. The gene fragment can be used to detect, amplify and/or isolate  
CC nicotianamine aminotransferase genes. 7 G;  
CC Sequence 23 BP; 3 A; 2 C; 4 T;  
SQ

| RESULT  | 3  |  |  |  |
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| AC      | V4 8148;   |  |  |  |
| DT      | 27-Oct-1998 (first entry)  |  |  |  |
| DE      | Nicotianamine aminotransferase 58148.62 molecular weight protein, gene |  |  |  |
| DS      | ; nicotianamine aminotransferase; plant; iron absorption;              |  |  |  |
| KW      | iron deficiency chlorosis.   |  |  |  |
| OS      | Gramineae sp.  |  |  |  |
| FH      | Location/qualifiers  |  |  |  |
| CDS     |  |  |  |  |
| FT      | 76..1731   |  |  |  |
| FT      | /*tag-   |  |  |  |
| FT      | product= "Nicotianamine transferase"                                   |  |  |  |

PN EF-000432 A2.  
PD 26-AUG-1998. 102891.  
PF 19-FEB-1998; 102891.  
PR 21-FEB-1997; JP-037499.  
PA (SUMO ) SUMITOMO CHEM CO LTD.  
PA Mori S, Nakaniishi H, Takahashi M;  
DR WPI: 98-439341/38.  
DR P-SPDB; W61643.  
PT New nicotianamine aminotransferase protein and DNA - useful for  
PT enhancing iron absorption of plant cells  
PS Claim 4: Page 14-15; 17pp; English.  
CC The nicotianamine aminotransferase can be used in a plasmid to transform  
CC plant cells to produce cells with enhanced iron absorption, and it is  
CC implied [though not stated] that plants with improved resistance to iron  
CC deficiency chlorosis in calcareous soils can be regenerated from the  
CC transformed cells. The gene fragment can be used to detect, amplify  
CC and/or isolate nicotianamine aminotransferase genes.  
SO Sequence 1910 bp. 462 A: 534 C: 546 G: 368 T;

| Query Match              | Best Local Similarity | Score 17.2;    | DB 1;          | Length 1910; | Gaps 0;   |
|--------------------------|-----------------------|----------------|----------------|--------------|-----------|
| Best Local Similarity    | 74.8%                 |                |                |              |           |
| Best Match <sub>16</sub> | 69.6%                 | Pred. No. 3.9; | 3. Missmatches | 4:           | Indels 0; |
| Conservative             |                       |                |                |              | Gaps 0;   |

|        |                              |                          |             |
|--------|------------------------------|--------------------------|-------------|
| Qy     | 1                            | gongtngartggaaattygcnnng | 23          |
|        |                              | :     :     :     :      |             |
| Db     | 460                          | GGGTGAGTGCATTCGCGGG      | 482         |
| RESULT | 4                            |                          |             |
| V26030 |                              |                          |             |
| ID     | V26030                       | standard; cDNA;          | 3258 BP.    |
| AC     | V26030;                      |                          |             |
| DT     | 28-AUG-1998                  | (first entry)            |             |
| DE     | Human transmembrane receptor | potential protein        | Htrp3 cDNA. |

|     |   |
|-----|---|
| KW  | Htrp3; transient receptor potential; trp protein; human;            |
| KW  | capacitative calcium ion entry; CCE; asthma; hypertension;          |
| KW  | diabetes; osteoporosis; osteogenesis; thrombosis; immunodeficiency; |
| KW  | gene therapy; ss.   |
| KW  | Homo sapiens.   |
| Key |   |
| CDS |   |
| FT  | Location/Qualifiers   |
| FT  | 291..3080   |
| FT  | /*tag= a  |
| PN  | W0980979-A1.  |
| PD  | 05-MAR-1998.  |
| PR  | 29-AUG-1997; U15247.  |
| PR  | 15-OCT-1996; US-729955.   |
| PR  | 30-AUG-1996; US-025111.   |
| PA  | (REGC ) UNIV CALIFORNIA.  |

1 Burdenet L, and A.  
2 DR WPI; 98-230369/20.  
3 DR WPI; 98-230369/20.  
4 P-PSDB: W55961.  
PT Controlling capacitative calcium ion entry into mammalian cells - by  
PT changing activity of transient receptor potential proteins, e.g. for  
PT treating asthma, hypertension etc.  
PT Claim 12: Page 31-35: 60pp; English.  
PS This cDNA clone codes for a human transient receptor potential (TRP) protein (see W55961), designated TRIP3, that is an essential part of the capacitative calcium ion entry (CCE) mechanism in human cells. The cDNA was isolated by subjecting human embryo cell line 293 CDR to RACE-PCR using primers (see V26035-38) based on the sequence of Genbank expressed sequence tag EST R34716. TRIP1 (see V26029) cDNA has also been isolated. CCE into a mammalian cell expressing a TRP protein required for CCE is controlled in a claimed method by treating the cell with an agent that increases or decreases the amount of biologically active TRP protein from its normal level. Agents that inhibit CCE are potentially useful for treating asthma, hypertension and osteoporosis, also for antithrombotic therapy, while those that stimulate CCE are used to treat type II diabetes and to induce bone formation. Primary immunodeficiency, if associated with TRP gene mutations, may be treated by gene therapy.

5 840 T;  
6 Sequence 3258 BP; 911 A;  
7 720 C;  
8 787 G;

OS Homo sapiens.  
PN WO980897-A1.  
PD 05-MAR-1998.  
PF 29-AUG-1997; U15247.  
PR 15-OCT-1996; US-72955.  
PR 30-AUG-1996; US-025111.  
PA (REGC) UNIV CALIFORNIA.  
PI Birnbaumer L, Zhu X;  
DR WPI: 98-230269/20.  
PT Controlling capacitative calcium ion entry into mammalian cells - by  
PT changing activity of transient receptor potential proteins, e.g. for  
PT treating asthma, hypertension etc.  
Disclosure: Page 9; 60pp; English.  
PS

PCR primers S1, S2, A1 and A2 (see V26035-38) are based on the expressed sequence tag R3416 from the GenBank dbEST database. They were used in the isolation and identification of Htrpc CDNA. Primary RACE-PCR amplifications of human embryonic kidney (HEK) 293 cell cDNA was performed using adaptor primer AP1 in combination with primer S1 for 3' amplification or A1 for 5' amplification. Nested PCR amplifications were performed using internal primers AP2 plus S2 for 3'RACE or AP2 plus A2 for 5'RACE. Positive clones were identified using oligonucleotides A1 and S2 for the 3', and 5', RACE analysis (see V26030) was obtained. It codes for a transient receptor potential (trp) protein (see W59961) that is essential for calcium ion entry into cells. The invention relates to methods for identifying cells with agents that raise or lower the amount of trp protein and which thereby control capacitative calcium ion entry into the cell, and for using trp proteins as targets for identifying such agents. Sequence 29 BP; 8 A; 10 C; 3 G; 8 T;

```

Query Match      67.8%; Score 15.6; DB 1; Length 29;
Best Local Similarity 65.2%; Pred. No. 14;
Matches 15; Conservative 3; Mismatches 5; Indels 0;
Gaps 0;
QY 1 ggcgtngartggaaayttgcnmg 23
Db 23 GATCTAGATGGAACTTGCCTCG 1

```

RESULT 6  
 V52248 standard; DNA; 11303 BP.  
 ID V52248  
 AC V52248  
 DT 23-OCT-1998 (first entry)  
 DE Streptococcus pneumoniae genome fragment SEQ ID NO:115  
 KW Streptococcus pneumoniae; S. pneumoniae; diagnosis; assay;  
 KW computer readable medium; vaccine; pharmaceutical composition; ds.  
 OS Streptococcus pneumoniae.

UD U-MAT-1296; U19588.  
 PF 30-OCT-1997; U19588.  
 PR 31-OCT-1996; US-029960.  
 PA (HUMA-) HUMAN GENOME SCI INC.  
 PT Barash SC, Choi GH, Dillon BJ, Dougherty BA, Fannon M,  
 PI Kunsch CA, Rosen CA;  
 WPI 98-27225/24.  
 DR Computer-readable medium with recorded Streptococcus pneumoniae  
 polynucleotide sequences - useful in diagnostic kits and assays, and  
 pharmaceutical compositions and vaccines for Streptococcus  
 pneumoniae.

The present invention describes a computer readable medium which has the nucleotide sequences SEQ ID NO:1 to 391 (V52134 to V5224) recorded on it, or a representative fragment or a sequence at least 95% identical to SEQ ID NO:1 to 391. The nucleotide sequences depicted in SEQ ID NO:1 to 391 (V52134 to V5224) are genomic fragments from Streptococcus pneumoniae. The present invention also describes an isolated nucleic acid molecule encoding a homologue of any of the fragments of the S. pneumoniae genome (SEQ ID NO:1 to 391) where the nucleic acid molecule is produced by a process comprising: (a) screening a genomic DNA library using as a probe a target sequence defined by any of the sequences in SEQ ID NO:1 to 391, identifying members of the library which contain sequences that hybridise to the target sequence and isolating the nucleic acid molecules from the members; or (b) isolating mRNA, DNA or cDNA produced from an organism, amplifying nucleic acid molecules whose nucleotide sequence is homologous to amplification primers derived from the fragment of the S. pneumoniae genome to prime the amplification and isolating the amplified sequences. The computer readable medium can be used in a computer-based system for identifying fragments of the S. pneumoniae genome of commercial importance, or expression modulating fragments of the S. pneumoniae genome. Products from the present invention can be used in diagnosis kits and assays, and pharmaceutical compositions and vaccines for S. pneumoniae.

Sequence 111303 BP: 3733 A: 1558 G: 3746 T:  
SQ 2266 Ci: 3733 A: 1558 G: 3746 T:

KW non-insulin dependent diabetes mellitus; diagnosis; human; ds.

OS Homo sapiens. Location/Qualifiers  
FH Key 184. .1122

FT CDS /\*tag= a /product= "human UCP3"  
FT FT 1..1032

FT /\*tag= b /note= "reading frame 1; contains internal stop codons which are not indicated in the corresponding protein (W81592)"

FT CDS /\*tag= c /note= "reading frame 2; contains internal stop codons which are not indicated in the corresponding protein (W81593)"

FT FT 2..1033

FT /\*tag= d /note= "reading frame 3; contains internal stop codons which are not indicated in the corresponding protein (W81594)"

FT CDS /\*tag= e /note= "reading frame 2; contains internal stop codons which are not indicated in the corresponding protein (W81589)"

FT FT 3..1032

FT /\*tag= f /note= "reading frame 3; contains internal stop codons which are not indicated in the corresponding protein (W81590)"

FT WO9845438-A1.  
PN PD 15-OCT-1998; US-892745.

PD 15-OCT-1998; US-06959.  
PF 08-APR-1998; US-06959.

PF 08-APR-1998; US-06959.

PR 09-JUL-1997; US-893447.  
PR 12-MAY-1997; US-046254.

PR 09-JUL-1997; US-046254.  
PR 12-MAY-1997; US-046254.

PI Flier JS; Lowell BB;  
DR WPI; 98-59483/50.  
PT New isolated uncoupling protein, UCP-3 - used to develop products for modulating thermogenesis in tissues, e.g. for treating obesity or muscle wasting caused by infection or cancer

CC This DNA encodes a human uncoupling protein 3 short form (UCP3sh) genes ('V7110 and V7112) encoding UCP3 proteins (W81587 and W81595) respectively. A host cell transformed with a construct comprising the UCP3 nucleic acid can be used for the recombinant production of the protein. The UCP3 is involved in the regulation of thermogenesis in mammals. The nucleic acids ('V7110 to V7112) can be used for identifying compounds which alter UCP3 activity. Enhancers of UCP3 can be used for enhancing protein catabolism in a mammal and can be used as anti-obesity drugs. Inhibitors of UCP3 can be used for inhibiting protein catabolism in a mammal such as inhibiting muscle wasting. They can be used for curtailing muscle wasting due to infection (e.g. HIV), cancer, tumour cachexia, muscle diseases (e.g. muscular dystrophy) or as a possible treatment for non-insulin dependent diabetes mellitus. The products can also be used for detection and diagnosis. The products can also be used for detection and diagnosis. Sequence 1220 BP; 267 A; 376 C; 345 G; 232 T;

PS Claim 3; Fig 1A-C; 98pp; English.

CC This DNA encodes a human uncoupling protein 3 (UCP3). A host cell transformed with a construct comprising the UCP3 nucleic acid can be used for the recombinant production of the protein. The UCP3 is involved in the regulation of thermogenesis in mammals. The nucleic acids ('V7110 to V7112) can be used for identifying compounds which alter UCP3 activity. Enhancers of UCP3 can be used for enhancing protein catabolism in a mammal and can be used as anti-obesity drugs. Inhibitors of UCP3 can be used for inhibiting protein catabolism in a mammal such as inhibiting muscle wasting. They can be used for curtailing muscle wasting due to infection (e.g. HIV), cancer, tumour cachexia, muscle diseases (e.g. muscular dystrophy) or as a possible treatment for non-insulin dependent diabetes mellitus. The products can also be used for detection and diagnosis. Sequence 1220 BP; 267 A; 376 C; 345 G; 232 T;

CC Best Local Similarity 64.3%; Score 14.8; DB 1; Length 1220; Pred. No. 66; Mismatches 4; Indels 0; Gaps 0;

Qy 2 cngtngartggayttgcnn 22  
Db 715 CTGTGAAAGGAACTTGCC 735

RESULT 9  
V71711 standard DNA; 1033 BP.  
AC V71711;  
DT 09-FEB-1999 (first entry)  
Human uncoupling protein 3 short form (UCP3sh) encoding DNA.  
Uncoupling protein 3; UCP; thermogenesis; mammal; enhancer; drug; protein catabolism; anti-obesity; inhibitor; muscle wasting; infection; HIV; cancer; tumour cachexia; muscle disease; diagnosis; human; ds.

KW non-insulin dependent diabetes mellitus; diagnosis; human; ds.

OS Homo sapiens. Location/Qualifiers  
FH Key 181. .1008  
FT CDS /\*tag= a /product= "UCP3L"  
FT

Query Match 64.3%; Score 14.8; DB 1; Length 1033;  
Best Local Similarity 61.9%; Pred. No. 64; Mismatches 4; Indels 0; Gaps 0;

Matches 13; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 2 cngtngartggayttgcnn 22  
DB 712 CTGTTGGAAAGGAACTTGCC 732

RESULT 10  
V72690 standard; cDNA; 1231 BP.  
ID V72690;  
AC V72690;  
DT 22-FEB-1999 (first entry)  
Human uncoupling protein UCP1 encoding cDNA.  
Human uncoupling protein; mitochondrial; gene therapy;  
adenosine triphosphate; mitochondria; skeletal muscle; gene therapy;  
thermogenesis; heart; obesity; cachexia; type II diabetes; hypertension;  
insulin sensitivity; neuromuscular disease; ss.  
Homo sapiens.  
Key  
FT  
CDS /\*tag= a /product= "UCP3L"  
FT

PN WO9850542-A1.  
 PD 12-NOV-1998.  
 PF 05-MAY-1998; E02645.  
 PR 07-MAY-1997; CH-001072.  
 PA (NOVS ) NOVARTIS-EREFINDUNGEN VERW GES MBH.  
 PI Boss O, Giacobino J, Muzzin P;  
 DR WPI; 98-610382/51.  
 P-PSDB; W83379.  
 PT New human skeletal muscle uncoupling proteins UCP3L and UCP3S - useful for controlling thermogenesis in human skeletal muscle and heart, e.g. for treating obesity and cachexia  
 Claim 1; Page 14-15; 26PP; English.  
 PS The present sequence encodes human uncoupling protein UCP3L. UCP3 uncouples oxidative phosphorylation and synthesis of adenosine triphosphate in the mitochondria of skeletal muscle. The coding sequences for UCP3L and UCP3S are useful for gene therapy of dysfunctions of thermogenesis in human skeletal muscle and heart which result from a lack of UCP3 and which can induce disorders such as obesity or cachexia. Antisense oligonucleotides to UCP3L and UCP3S can be used for correcting an excess of UCP3. Modification of endogenous UCP3 activity (using activators or inhibitors of UCP3) is used to induce bodyweight loss (loss of adipose mass and maintenance of the lean mass) in all types of obesity by promoting the dissipation of energy; for preventing an excessive weight regain following a restrictive food diet or after ceasing a physical training programme; for preventing and treating type II diabetes by improving sensitivity to insulin; for preventing hypertension, for increasing muscle mass in states of cachexia; for treatment of insufficiencies or disturbances of cardiac rhythm due to a dysfunction of UCP3; and for the treatment of neuromuscular diseases due to a dysfunction of UCP3, and for the treatment of neuromuscular diseases due to a dysfunction of UCP3. The uncoupling proteins can also be used to raise antibodies, e.g. for diagnosis. Knowledge of the UCP3 genes allows generation of transgenic animals, e.g. for screening substances which modify UCP3 expression or activity or for investigating the biological role of UCP3.  
 Sequence 1231 BP; 271 A; 371 C; 345 G; 244 T; 215 T; 321 G; 245 A; 215 T;

PT heart, e.g. for treating obesity and cachexia  
 Claim 3; Page 17-18; 26PP; English.

PS The present sequence encodes human uncoupling protein UCP3S. UCP3 uncouples oxidative phosphorylation and synthesis of adenosine triphosphate in the mitochondria of skeletal muscle. The coding sequences for UCP3L and UCP3S are useful for gene therapy of dysfunctions of thermogenesis in human skeletal muscle and heart which result from a lack of UCP3 and which can induce disorders such as obesity or cachexia. Antisense oligonucleotides to UCP3L and UCP3S can be used for correcting an excess of UCP3. Modification of endogenous UCP3 activity (using activators or inhibitors of UCP3) is used to induce bodyweight loss (loss of adipose mass and maintenance of the lean mass) in all types of obesity by promoting the dissipation of energy; for preventing an excessive weight regain following a restrictive food diet or after ceasing a physical training programme; for preventing and treating type II diabetes by improving sensitivity to insulin; for preventing hypertension, for increasing muscle mass in states of cachexia; for treatment of insufficiencies or disturbances of cardiac rhythm due to a dysfunction of UCP3; and for the treatment of neuromuscular diseases due to a dysfunction of UCP3, and for the treatment of neuromuscular diseases due to a dysfunction of UCP3. The uncoupling proteins can also be used to raise antibodies, e.g. for diagnosis. Knowledge of the UCP3 genes allows generation of transgenic animals, e.g. for screening substances which modify UCP3 expression or activity or for investigating the biological role of UCP3.

Query Match Score 64.3%; Score 14.8%; DB 1; Length 1132;  
 Best Local Similarity 61.9%; Pred. No. 65; Gaps 0;  
 Matches 13; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 2 cngtngartggaaattygcnm 22  
 DB 685 CTGTCGAAGGACTTGGCC 705

RESULT 12  
 ID V84254 standard; cDNA; 2340 BP.  
 AC W84254.  
 DT 12-APR-1999 (first entry)  
 DE Human uncoupling protein 3 (UCP3) cDNA.  
 KW Uncoupling protein 3; UCP3; human; obesity; diabetes;  
 KW hyperinsulinaemia; hypermetabolism; gene therapy; ds.  
 OS Homo sapiens.  
 PH Location/Qualifiers  
 FT Key  
 FT CDS 344..1282  
 FT /\*tag= ^a  
 FT /\*tag= ^a  
 FT /product= "UCP3S"  
 FT /note= "this region is specifically claimed in  
 Claim 2"

Query Match Score 64.3%; Score 14.8%; DB 1; Length 1231;  
 Best Local Similarity 61.9%; Pred. No. 66;  
 Matches 13; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 2 cngtngartggaaattygcnm 22  
 DB 685 CTGTCGAAGGACTTGGCC 705

RESULT 11  
 ID V72691 standard; cDNA; 1132 BP.  
 AC V72691.  
 DT 22-FEB-1999 (first entry)  
 DE Human uncoupling protein UCP3S encoding cDNA.  
 KW Human; uncoupling protein; UCP3L; UCP3S; oxidative phosphorylation; adenosine triphosphate; mitochondria; skeletal muscle; gene therapy; thermogenesis; heart; obesity; cachexia; type II diabetes; hypertension; insulin sensitivity; neuromuscular disease; ss.  
 KW Homo sapiens.  
 OS Location/Qualifiers  
 FT 154..991  
 FT /\*tag= ^a  
 FT /product= "UCP3S"  
 PN WO9850542-A1.  
 PD 12-NOV-1998.  
 PF 05-MAY-1998; E02645.  
 PR 07-MAY-1997; CH-001072.  
 PA (NOVS ) NOVARTIS AG.  
 PA (NOVS ) NOVARTIS-BREFINDUNGEN VERW GES MBH.  
 PI Boss O, Giacobino J, Muzzin P;  
 DR WPI; 98-110382/51.  
 DR P-PSDB; W83380.  
 PT New human skeletal muscle uncoupling proteins UCP3L and UCP3S - useful for controlling thermogenesis in human skeletal muscle and

PT heart, e.g. for treating obesity and cachexia  
 Claim 3; Page 17-18; 26PP; English.

PS The present sequence encodes human uncoupling protein 3 (UCP3, see W88279), a novel protein that is involved in energy expenditure and body weight regulation and whose expression is mostly limited to skeletal muscle. The clone was isolated from a foetal brain cDNA library using primers (see W84259-64) based on isolated EST clones (see W84255-58) and Bluescript vector sequences. The invention additionally provides related recombinant expression vectors, recombinant host cells and purified forms of the UCP3 protein. The UCP3 polypeptides and transformed recombinant cell lines can

CC be used for identifying modulators of UCP3 activity. Such modulators can be used for treating diseases such as obesity and diabetes, by manipulating the interrelated process of balancing food intake, energy expenditure and glucose metabolism within the patient. They can also be used to treat hyperactive conditions of energy expenditure which originate in the mitochondria of skeletal muscle. UCP3 nucleic acids are useful in gene therapy of obesity and obesity-related indications, including diabetes, and of mitochondrial-associated hypermetabolism.

CC mitochrondrial-associated hypermetabolism. Sequence - 2340 BP; 606 A; 638 C; 633 G; 463 T;

Query Match Score 14.8; DB 1; Length 2340;

Best Local Similarity 61.9%; Pred. No. 73; Mismatches 4; Indels 0; Gaps 0;

Matches 13; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

Qy 2 crngtngartggaaatyygcnn 22

Dy 1 |||:|||:|||:|||:|||:|||:

Db 875 CTGTGGAAAGGAACTTGCC 895

RESULT 13

ID N92379/c standard; DNA; 3094 BP.

AC N92379; 1992 (first entry)

DT 15-MAR-1992 Sequence of the 1.7kb cDNA molecule encoding antibodies 7D1,7D4

DE and 20C6.

KW Vaccine; poultry; bivalent vaccine; ss.

OS Eimeria.

Location/Qualifiers

FT 9\_3002

PT EP-344808-A.

PY 06-DEC-1989; 110056;

PN 02-JUN-1988; US-2027-21.

PA (HOFF ) HOFFMANN-LA ROCHE AG.

PI Altenburger W, Binger MH, Chizzonite RA, Kramer RA,

PT Lomedico PT, McAndrew SJ;

DR WPI: 89-358220/49.

P-PSDB; P93706.

PT DNA sequences encoding Eimeria surface antigens - used in poxvirus vectors as a vaccine to protect chicks against coccidiosis.

PT Claim 10; Fig 20A-D; 78pp; English.

CC The inventors claim a new protein which comprises one or more immunoreactive and/or antigenic determinants of an Eimeria surface antigen of mol. wt. 28,37,120 or more than 200 kD which specifically binds to one or more monoclonal antibody (mAb) from ATCC HB 9707-9712 (see P93703-6). Also new are DNA encoding the protein (see N92516-9), and a vaccine comprising one or more proteins. Vaccine utility can be enhanced by inserting additional genes into the carrier virus (see P91652).

CC Sequence 3094 BP; 617 A; 834 C; 846 G; 796 T;

SQ OS Eimeria tenella.

KW sporozoite; ss; Eimeria tenella. Location/Qualifiers

FT Key

FT CDS 7\_2997

FT /\*tag= a

FT US5661015-A.

FT PD 03-JUN-1988; 202721.

FT PR 20-DEC-1991; US-812349.

FT PR 03-JUN-1988; US-202721.

FT PA (HOFF ) HOFFMANN LA ROCHE INC.

PI Altenburger W, Binger M, Chizzonite RA, Kramer RA,

PT Lomedico PT, McAndrew SJ;

DR WPI: 97-434379/40.

P-PSDB; W33624.

PT New DNA from Eimeria tenella and related immunogenic polypeptides -

PT useful in vaccines to protect poultry against coccidiosis.

PS Claim 2; FIG 33A-B; 72pp; English.

CC This cDNA sequence comprises the coding sequence for a 45 kDa protein (see W33624) that is recognised by monoclonal antibody 7B2 (ATCC HB 9712). This antibody also specifically reacts with an Eimeria tenella 200 kDa surface antigen that is present in the sporozoite developmental stage. The clone (see also T93596) was obtained from a cDNA library by immunological screening with conventional antibodies raised against Eimeria antigens. The invention provides DNA sequences (see W31582-84 and W33621-26), recombinant vectors containing such DNA sequences, transformed microorganisms containing such vectors, and methods for producing the antigens using the transformed microorganisms. Methods are also provided for protecting poultry against coccidiosis using the Eimeria surface antigens. The surface antigens are administered either as purified proteins or in the form of DNA encoding the proteins in a viral vector such as a vaccinia virus. The vaccines may produce antibodies that are cross-reactive with other Eimeria species.

CC Sequence 2997 BP; 583 A; 815 C; 769 T;

SQ DR WPI: 97-434379/40.

Query Match Score 14.6; DB 1; Length 2997;

Best Local Similarity 63.6%; Pred. No. 97; Matches 14; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

Qy 1 gcngtngartggaaatyygcnn 22

Db 254 GCAGGGGAGTGGAGTTCGCGA 233

RESULT 15

ID T93596/c standard; cDNA; 3094 BP.

AC T93596; 1992 (first entry)

DT 21-MAY-1998

DE Eimeria tenella sporozoite 45 kDa surface antigen cDNA.

KW Coccidiosis; vaccine; poultry; protozoan; parasite; antigen; sporozoite; ss.

OS Eimeria tenella.

Location/Qualifiers

FT 9\_3005

FT /\*tag= a

FT /transl\_except= (pos:600..602, aa:Tyr)

FT /transl\_except= (pos:603..605, aa:Ser)

FT /transl\_except= (pos:606..608, aa:Gly)

FT /transl\_except= (pos:612..614, aa:Ala)

FT /transl\_except= (pos:615..617, aa:Leu)

FT misc\_difference 2001

FT /\*tag= b

FT /note= "base 2001 is given as '8' in Fig20 of the specification"

RESULT 14

ID T93597/c standard; cDNA; 2997 BP.

AC T93597; 1992 (first entry)

DT 21-MAY-1998

DE Eimeria tenella sporozoite 45 kDa surface antigen cDNA.

KW Coccidiosis; vaccine; poultry; protozoan; parasite; antigen;

PR 03-JUN-1988; US-202721.  
 PA (HOFF ) HOFFMANN LA ROCHE INC.  
 PI Altenburger W, Binger M, Chizzonite RA, Kramer RA,  
 PI Lomedico PT, McAndrew SJ,  
 DR WPI; 97-34379/40.  
 DR P-PSDB; W33621.  
 PT New DNA from *Emeria tenella* and related immunogenic polypeptides -  
 PT useful in vaccines to protect poultry against coccidiosis  
 PS Claim 1; Fig 20A-D; 72pc; English.  
 CC This cDNA clone includes a coding region for a 45 kDa protein (see  
 CC W33621) that is recognised by monoclonal antibody 7B2 (ATCC HB 9112).  
 CC This antibody also specifically reacts with an *Emeria tenella* 200  
 CC kDa surface antigen that is present in the sporozoite developmental  
 CC stage. The clone was obtained from a cDNA library by immunological  
 CC screening with monoclonal antibodies raised against *Emeria*  
 CC antigens. The first and last 7 nucleotides of the sequence are  
 CC derived from linker sequences used in the cloning procedure. The  
 CC invention provides DNA sequences (see T93503-98) coding for *Emeria*  
 CC surface antigens (see W31582-84 and W33621-26), recombinant vectors  
 CC containing such DNA sequences, transformed microorganisms  
 CC containing such vectors, and methods for producing the antigens  
 CC using the transformed microorganisms. Methods are also provided  
 CC for protecting poultry against coccidiosis using the *Emeria*  
 CC surface antigens. The surface antigens are administered either as  
 CC purified proteins or in the form of DNA encoding the proteins in  
 CC a viral vector such as a vaccinia virus. The vaccines may produce  
 CC antibodies that are cross-reactive with other *Emeria* species.  
 Sequence 3094 BP; 619 A; 844 G; 796 T;  
 SQ

Query Match 63.5%; Score 14.6; DB 1; Length 3094;  
 Best Local Similarity 63.6%; Pred. No. 97;  
 Matches 14; Conservative 3; Mismatches 5; Indels 0; Caps 0;  
 Qy 1 gcnctgatggatggaaatyytgcnn 22  
 Db 262 GCAGGGAGGTTGGAAAGTTCSCGA 241

Search completed: October 1, 1999, 15:36:18  
 Job time: 6116 sec



GenCore version 4.5  
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OM nucleic - nucleic search, using sw model  
Run on: October 1, 1999, 15:03:35 ; Search time 1096.08 Seconds  
(without alignments)  
41.391 Million cell updates/sec

Title: US-09-026-400-5  
Perfect score:  
23

Sequence:  
1 gngtgartggaaatygcnmng 23

Scoring table: IDENTITY\_NUC

Searched: 2546578 seqs., 986266752 residues

Database : EST:\*

| Result No. | Score | Query Match Length | DB ID              | Description        |
|------------|-------|--------------------|--------------------|--------------------|
| C 1        | 16.8  | 73 .0              | AA660414           | AA660414 00290 Mtr |
| C 2        | 15.8  | 68 .7              | H1322 Ym13a03.s1   |                    |
| C 3        | 15.8  | 68 .7              | H13911 Yj08a05.r1  |                    |
| C 4        | 15.8  | 68 .7              | H56483 Yt87a1.s1   |                    |
| C 5        | 15.8  | 68 .7              | H9744 Yx12c12.s1   |                    |
| C 6        | 15.8  | 68 .7              | W0531 mb47b04.r1   |                    |
| C 7        | 15.8  | 68 .7              | W41942 m64e07.r1   |                    |
| C 8        | 15.8  | 68 .7              | W66228 zd29g09.s1  |                    |
| C 9        | 15.8  | 68 .7              | W76544 zd6a09.s1   |                    |
| C 10       | 15.8  | 68 .7              | AA03148 ms51901.r  |                    |
| C 11       | 15.8  | 68 .7              | AA01075 ze2ff01.r  |                    |
| C 12       | 15.8  | 68 .7              | AA036822 zk29b05.r |                    |
| C 13       | 15.8  | 68 .7              | AA038824 m195g07.r |                    |
| C 14       | 15.8  | 68 .7              | AA05760 zl193d09.s |                    |
| C 15       | 15.8  | 68 .7              | W82322 mf04911.r1  |                    |
| C 16       | 15.8  | 68 .7              | W82575 mf04a11.r1  |                    |
| C 17       | 15.8  | 68 .7              | AA059678 mj75d04.r |                    |
| C 18       | 15.8  | 68 .7              | AA060109 m171g12.r |                    |
| C 19       | 15.8  | 68 .7              | AA061890 m192h08.r |                    |
| C 20       | 15.8  | 68 .7              | AA075167 mm89g04.s |                    |
| C 21       | 15.8  | 68 .7              | AA082814 zn25b04.r |                    |
| C 22       | 15.8  | 68 .7              | AA103480 mo24h04.r |                    |
| C 23       | 15.8  | 68 .7              | AA132788 zo22d08.s |                    |
| C 24       | 15.8  | 68 .7              | AA145601 mr63f01.r |                    |
| C 25       | 15.8  | 68 .7              | AA147813 zo48b09.s |                    |
| C 26       | 15.8  | 68 .7              | AA151511 zl26h07.r |                    |
| C 27       | 15.8  | 68 .7              | AA183476 mo97f03.r |                    |
| C 28       | 15.8  | 68 .7              | AA187994 ZP67h06.s |                    |
| C 29       | 15.8  | 68 .7              | AA206634 ZQ80C1.s  |                    |
| C 30       | 15.8  | 68 .7              | AA225589 mv70b12.r |                    |
| C 31       | 15.8  | 68 .7              | AA243010 zr25h03.s |                    |
| C 32       | 15.8  | 68 .7              | AA250948 zs06h11.r |                    |
| C 33       | 15.8  | 68 .7              | AA266452 vaoa06.r  |                    |
| C 34       | 15.8  | 68 .7              | AA336513 EST41591  |                    |
| C 35       | 15.8  | 68 .7              | AA447651 zw97g04.s |                    |
| C 36       | 15.8  | 68 .7              | AA472282 v01b03.r  |                    |
| C 37       | 15.8  | 68 .7              | AA474481 vd55e09.r |                    |
| C 38       | 15.8  | 68 .7              | AA475447 vh15a11.r |                    |
| C 39       | 15.8  | 68 .7              | AA516883 vh88h03.r |                    |
| C 40       | 15.8  | 68 .7              | AA522284 aa75h07.s |                    |
| C 41       | 15.8  | 68 .7              | AA555952 vi64g01.r |                    |
| C 42       | 15.8  | 68 .7              | AA594843 no21b08.s |                    |
| C 43       | 15.8  | 68 .7              | AA610937 nr72e09.s |                    |
| C 44       | 15.8  | 68 .7              | AA662313 nu97q03.s |                    |
| C 45       | 15.8  | 68 .7              | AI540076 td09a05.x |                    |

## SUMMARIES

RESULT 1  
AA660414/c  
LOCUS AA660414  
DEFINITION 00290 MtrHE Medicago truncatula cDNA 5', mRNA sequence.  
NID 92604458  
VERSION AA660414.1  
KEYWORDS EST

EST 10-NOV-1997  
mRNA  
mtrHE Medicago truncatula cDNA 5', mRNA sequence.

## ALIGNMENTS

SOURCE ORGANISM Medicago truncatula; Streptophyta; Embryophyta; Tracheophyta; Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; eudicotyledons; core eudicots; Rosidae; eurosids I; Fabales; Fabaceae; Papilionoideae; Medicago.

REFERENCE Covit,P.A., Smith,L.S. and Long,S.R.

AUTHORS Unpublished sequence tags from a root hair-enriched *Medicago truncatula* cDNA library

TITLE Unpublished (1997)

JOURNAL On Sep 12, 1996 this sequence version replaced gi:1290520.

COMMENT Contact: Long SR  
Department of Biological Sciences and Howard Hughes Medical Institute  
Stanford University  
Gibbert Biology, Stanford, CA 94305-5020, USA  
Tel: 650 723 3232  
Fax: 650 725 8309  
Email: fa.srl@forsythe.stanford.edu

FEATURES source

Location/Qualifiers Seq primer T3.

Source /cultivar="Medicago truncatula"  
/cultivar="Jemalong"  
/db\_xref="taxon:3880"  
/clone\_lib="MRHE"

Tissue\_type="Root hairs & tips"  
dev\_stage="2-3 day old seedlings"  
note="Organ: Root; Vector: PBK-CMV; Site\_1: ECORI;  
Site 2: XbaI; cDNA was synthesized from a pooled mRNA prep  
from elongating root hairs (30% w/w) and 2-3cm root tips  
(70% w/w). XbaI-oligo-dT linker primer and ECORI  
adaptors were used. cDNAs was cloned unidirectionally  
into lambda ZAP Express (Stratagene), amplified, and  
mass-excised into PBK-CMV vector plasmids. More  
information is available at <http://bio.SRL.stanford.edu>."

BASE COUNT 187 a 149 c 133 g 178 t 46 others

ORIGIN

Query Match 73.0%; Score 16.8%; DB 36; Length 693;  
Best Local Similarity 65.2%; Pred. No. 38; Mismatches 4; Indels 0; Gaps 0;

Matches 15; Conservative 4; Mismatches 3

Qy 1 gcnctgatggaaatyygcngm 23  
| | | |:| | |:||:||:|:  
Db 206 GCAGTGAGTGAAATTGGACG 184

RESULT 2

H11322 LOCUS H11322 539 bp mRNA EST 26-JUN-1995  
DEFINITION yml3a03\_s1 Soares infant brain INIB Homo sapiens cDNA clone IMAGE:47320 3', mRNA sequence.

ACCESSION H11322  
NID 9876142  
VERSION H11322.1 GI:876142  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 539)  
Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J., Raskin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston,R., Williamson,A., Wohldmann,P. and Wilson,R.

AUTHORS The WashU-Merck EST Project  
Unpublished (1995)

TITLE The WashU-Merck EST Project  
COMMENT On May 10, 1995 this sequence version replaced gi:805432.

Contract: wilson RK  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: est@watson.wustl.edu  
Insert Size: 1450  
High quality sequence stops: 410  
Source: IMAGE Consortium, LNL  
This clone is available royalty-free through LNL ; contact the  
IMAGE Consortium (info@image.llnl.gov) for further information.  
Insert Length: 1450 Std Error: 0.00  
Seq Primer: Promega -21ml3  
High quality sequence stop: 410.  
Location/Qualifiers

FEATURES source

Location/Qualifiers Seq primer T3.

Source /organism="Homo sapiens"  
/db\_xref="SDB:420061"  
/db\_xref="taxon:9606"  
/clone="IMAGE:47520"  
/lab\_host="DH10B (ampicillin resistant)"  
/note="Organ: whole brain; Vector: Lambda BA; Site\_1: Not I; Site\_2: Hind III; 1st strand cDNA was primed with a Not I - Oligo(dT) primer [5', AACTGGAGAATCGGCCGAGGATTTTTTTTTTTTTTTTT 3']; double-stranded cDNA was ligated to Hind III adaptors (Pharmacia), digested with Not I and directionally cloned into the Not I and Hind III sites of the Lafmid BA vector. Library went through one round of normalization. Library constructed by Bento Soares and M.Fabima Bonaldo."

BASE COUNT 135 a 132 c 131 g 135 t 6 others

ORIGIN

Query Match 66.7%; Score 15.8%; DB 22; Length 539;  
Best Local Similarity 63.6%; Pred. No. 1.3e+02; Mismatches 4; Indels 0; Gaps 0;

Qy 2 cngtngatggaaatyygcngm 23  
| | | |:| | |:||:||:|:  
Db 453 CAGTGAGGGAACTTCGCAG 432

RESULT 3

H13911 LOCUS H13911 433 bp mRNA EST 27-JUN-1995  
DEFINITION Y108905 r1 Soares placenta Nb2HP Homo sapiens cDNA clone IMAGE:148184 5', mRNA sequence.

ACCESSION H13911  
NID 9878731  
VERSION H13911.1 GI:878731  
KEYWORDS EST.  
SOURCE human.  
ORGANISM Homo sapiens  
Eukaryota; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 433)  
Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J., Raskin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston,R., Williamson,R., Williamson,A., Wohldmann,P. and Wilson,R.

AUTHORS The WashU-Merck EST Project  
Unpublished (1995)

TITLE The WashU-Merck EST Project  
COMMENT Contact: wilson RK  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800

Fax: 314 286 1810  
 Email: est@watson.wustl.edu  
 Insert Size: 551  
 High quality sequence stops: 331  
 Source: IMAGE Consortium, LNL  
 This clone is available royalty-free through LNL ; contact the  
 IMAGE Consortium (info@image.lnl.gov) for further information.  
 Insert Length: 591 Std Error: 0.00  
 Seq. Primer: M13RP1  
 High quality sequence stop: 331.  
 Location/Qualifiers  
 1. .433  
 /organism="Homo sapiens"  
 /db\_xref="GDB:55920"  
 /db\_xref="Taxon:9606"  
 /clone="IMAGE:148184"  
 /clone\_id="Soares placenta Nb2HP"  
 /sex="Female"  
 /dev\_stage="Placenta obtained at birth (full term)"  
 /lab\_host="DH10B (ampicillin resistant)"  
 /note="Organ: Placenta; Vector: pT7T3D (Pharmacia) with a  
 modified polylinker. Site\_1: Not I; Site\_2: Eco RI; 1st  
 strand cDNA was primed with a Not I - oligo(dt) primer [5'  
 AACTGGAGAATTCGGCCAGCAATTCTTTTTTTTTTTTTTTTTTTTTTT  
 ], double-stranded cDNA was ligated with Eco RI adaptors  
 (Pharmacia), digested with Not I and cloned into the Not I  
 and Eco RI sites of the modified pT7T3 vector. Library  
 went through one round of normalization. Library  
 constructed by Bento Soares and M.Fatima Bonaldo. "  
 BASE COUNT  
 ORIGIN  
 103 a 105 c 120 g 103 t 2 others

RESULT 4  
 LOCUS H56483 281 bp mRNA  
 DEFINITION EST 02-OCT-1995 Homo sapiens cDNA clone  
 IMAGE:231310 3' similar to gb:X71973 PHOSPHOLIPID HYDROPEROXIDE  
 ACCESSION H56483  
 VERSION 91005127  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
 Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 281)  
 AUTHORS Hillier,L., Lennon,G., Becker,M., Bonaldo,M.F., Chiappelli,B.,  
 Chissoe,S., Dietrich,N., Dubroque,T., Favello,A., Gish,W.,  
 Hawkins,M., Hullman,M., Kucaba,T., Lacy,M., Le,N.,  
 Mardis,E., Moore,B., Morris,M., Parsons,J., Prange,C., Riffkin,L.,  
 Rohlfing,T., Schallenberg,K., Soares,M.B., Tan,F., Thierry-Mag,J.,  
 Trevassis,E., Underwood,K., Wohldmann,P., Waterston,R., Wilson,R.,  
 and Marra,M.  
 TITLE Generation and analysis of 280,000 human expressed sequence tags  
 JOURNAL Genome Res. 6 (9), 807-838 (1996)  
 MEDLINE 97044478  
 COMMENT On May 9, 1995 this sequence version replaced gi:803035.

Query Match 68.7%; Score 15.8%; DB 22; Length 433;  
 Best Local Similarity 63.6%; Prod. No. 1.2e+02; Indels 0; Gaps 0;  
 Matches 14; Conservative 4; Mismatches 04;

QY 2 cngtngartggaaatggcgtgcngm 23  
 Db 108 CAGTGAGGGAACTTGTCCAG 129

RESULT 5  
 LOCUS H98744 463 bp mRNA  
 DEFINITION EST 15-DEC-1995 Homo sapiens cDNA clone  
 IMAGE:261526 3' mRNA sequence.  
 ACCESSION H98744  
 VERSION 91123412  
 KEYWORDS EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
 Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 REFERENCE 1 (bases 1 to 463)  
 AUTHORS Hillier,L., Clark,N., Dubroque,T., Elliston,K., Hawkins,M.,  
 Holman,M., Hullman,M., Kucaba,T., Le,M., Lennon,G., Marra,M.,  
 Parsons,J., Rifkin,L., Rohlfing,T., Soares,M., Tan,F.,  
 Trevaskis,E., Waterston,R., Williamson,A., Wohldmann,P., and  
 Wilson,R.  
 TITLE The WashU-Merck EST Project  
 JOURNAL Unpublished (1995)  
 COMMENT On Nov 22, 1995 this sequence version replaced gi:1071096.

Contact: Wilson RK  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: est@watson.wustl.edu  
 High quality sequence stops: 377

Source: IMAGE Consortium, LINL  
 This clone is available royalty-free through LINL ; contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.  
 Insert Length: 1012 Std Error: 0.00  
 Seq Primer: m13 -40 forward  
 High quality sequence stop: 377.

FEATURES source

1. . 463  
 /organism="Homo sapiens"  
 /db\_xref="GDB:3871168"  
 /note="vector: pT73D (Pharmacia) with a modified  
 polylinker; Site\_1: Not I; Site\_2: Eco RI; 1st strand cDNA  
 was primed with a Not I - oligo(dt) primer [5',  
 TGTTCACATCTAAGTGAGGGCGCCATTTTTTTTTTTTTTTT 3'],  
 double-stranded cDNA was size selected, ligated to Eco RI  
 adaptors (Pharmacia), digested with Not I and cloned into  
 the Not I and Eco RI sites of a modified pT73 vector  
 (Pharmacia). Library went through one round of  
 normalization to a Cot = 5. Library constructed by Bento  
 Soares and M.Fatima Bonaldo. RNA was kindly provided by  
 Dr. Minoru Ko (Wayne State University)."

/sex="Male"  
 /tissue-type="melanocyte"  
 /lab\_host="DH10B (ampicillin resistant)"  
 /note="vector: pT73D (Pharmacia) with a modified  
 polylinker; Site\_1: Not I; Site\_2: Eco RI; 1st strand cDNA  
 was primed with a Not I - oligo(dt) primer [5',  
 TGTTCACATCTAAGTGAGGGCGCCATTTTTTTTTTTTTT 3'],  
 double-stranded cDNA was size selected, ligated to Eco RI  
 adaptors (Pharmacia), digested with Not I and cloned into  
 the Not I and Eco RI sites of a modified pT73 vector  
 (Pharmacia). Library constructed by Bento Soares and  
 M.Fatima Bonaldo. RNA from normal foreskin melanocytes  
 (PF534) was kindly provided by Dr. Anthony P. Albino."

BASE COUNT ORIGIN

Query Match Score 15.8; DB 24; Length 463;  
 Best Local Similarity 63.6%; Prod. No. 1.2e+03;  
 Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

RESULT 6  
 W08531 421 bp mRNA EST  
 LOCUS W08531  
 DEFINITION IMAGE:332527 5 , mRNA sequence.  
 ORGANISM Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
 Reference Eutheria; Rodentia; Sciurognathii; Murinae; Mus.  
 Authors Marr,M., Hillier,L., Allen,M., Dietrich,N., Dubuque,T.,  
 Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,  
 Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,  
 Theising,B., Wyllie,T., Lennon,G., Soares,B., Wilson,R. and  
 Waterston,R.  
 Title Unpublished (1996)  
 Journal The WashU-HMMI Mouse EST Project  
 Comment On Nov 29, 1993 this sequence version replaced gi:693673.

Contact: Marra M/Mouse EST Project  
 WashU-HMMI Mouse EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1800  
 Email: mouseest@wustl.edu  
 This clone is available royalty-free through LINL ; contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.

REFERENCE AUTHORS

Title JOURNAL COMMENT

On Nov 29, 1993 this sequence version replaced gi:430122.

Contact: Marra M/Mouse EST Project  
 WashU-HMMI Mouse EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1800  
 Email: mouseest@wustl.edu  
 This clone is available royalty-free through LINL ; contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.

Seq primer: ETPrimer

FEATURES source

High quality sequence stop: 403.  
 Location/Qualifiers  
 1. . 421  
 /organism="Mus musculus"  
 /db\_xref="taxon:10090"  
 /map="1"  
 /clone="IMAGE:332527"  
 /clone.lib="Soares mouse p3NMF19.5"  
 /dev\_stage="19.5 dpc total fetus"  
 /lab\_host="DH10B (ampicillin resistant)"  
 /note="vector: pT73D (Pharmacia) with a modified  
 polylinker; Site\_1: Not I; Site\_2: Eco RI; 1st strand cDNA  
 was primed with a Not I - oligo(dt) primer [5',  
 TGTTACCATCTAAGTGAGGGCGCCATTTTTTTTTTTTTTTT 3'],  
 double-stranded cDNA was size selected, ligated to Eco RI  
 adaptors (Pharmacia), digested with Not I and cloned into  
 the Not I and Eco RI sites of a modified pT73 vector  
 (Pharmacia). Library went through one round of  
 normalization to a Cot = 5. Library constructed by Bento  
 Soares and M.Fatima Bonaldo. RNA was kindly provided by  
 Dr. Minoru Ko (Wayne State University)."

BASE COUNT ORIGIN

Query Match Score 15.8; DB 25; Length 421;  
 Best Local Similarity 63.6%; Prod. No. 1.2e+02;  
 Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 2 cngtnqartggaaayttgycnmg 23  
 Db 104 CAGTTGGAAACTTGCAG 125

RESULT 7  
 W41942 528 bp mRNA EST  
 LOCUS mc64e07.r1 Soares mouse embryo NDME13.5 14.5 Mus musculus cDNA  
 DEFINITION clone IMAGE:333316 5 , mRNA sequence.

ACCESSION W41942  
 NID 91325656  
 VERSION W41942.1  
 KEYWORDS GI:1325656  
 SOURCE EST.  
 ORGANISM Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
 Reference Eutheria; Rodentia; Sciurognathii; Murinae; Mus.  
 Authors Marr,M., Hillier,L., Allen,M., Dietrich,N., Dubuque,T.,  
 Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,  
 Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,  
 Theising,B., Wyllie,T., Lennon,G., Soares,B., Wilson,R. and  
 Waterston,R.  
 Title Unpublished (1996)  
 Journal The WashU-HMMI Mouse EST Project  
 Comment On Apr 14, 1993 this sequence version replaced gi:693673.

Contact: Marra M/Mouse EST Project  
 WashU-HMMI Mouse EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1800  
 Email: mouseest@wustl.edu  
 This clone is available royalty-free through LINL ; contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.

REFERENCE AUTHORS

Title JOURNAL COMMENT

On Nov 29, 1993 this sequence version replaced gi:430122.

Contact: Marra M/Mouse EST Project  
 WashU-HMMI Mouse EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1800  
 Email: mouseest@wustl.edu  
 This clone is available royalty-free through LINL ; contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.

Seq primer: ETPrimer

|  |  |                        |               |             |
|--|--|------------------------|---------------|-------------|
| /db_xref="taxon:1090"  |  |                        |               |             |
| /clone="IMAGE:353316"  |  |                        |               |             |
| /clone_lib="Soares mouse embryo NbME13.5 14.5"   |  |                        |               |             |
| /sex="unknown"   |  |                        |               |             |
| /tissue_type="embryo"  |  |                        |               |             |
| /dev_stage="13.5-14.5dpc total fetus"  |  |                        |               |             |
| /lab_host="DHL0B"  |  |                        |               |             |
| /note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site:1: Not I; Site:2: Eco RI; 1st strand CDNA was primed with a Not I - Oligo(dT) primer [5'-TGTTACCAATTGAACTGGGACGGCGCAATTTTTTTTTTTTTT 3'], double-stranded cDNA was size selected, ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library went through one round of normalization to a Cot = 5. Library constructed by M. Fatima Bonaldo. This library was constructed from the same fetus as the fetal lung library, Soares fetal lung NbHH19W." |  |                        |               |             |
| 14.5dpc embryos (total RNA provided by Minoru Ko, Wayne State Univ., from 2 ); double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization, and was constructed by Bento Soares and M.Fatima Bonaldo."  |  |                        |               |             |
| BASE COUNT   | 129  | a                      | 134           | c           |
| ORIGIN   | 148  | g                      | 117           | t           |
| Query Match  | 68.7%  | Score 15.8;            | DB 26;        | Length 528; |
| Best Local Similarity  | 63.6%  | Pred. No. 1.e+02;      | Indels 0;     | Gaps 0;     |
| Matches 14;  | Conservative   | 4;                     | Mismatches 4; |             |
| Qy   | 2  | cngtgartgaaatygcnmg 23 |               |             |
| Db   | 141  | CAGTGGGGAACTTTCGAG 162 |               |             |
| RESULT   | 8  |                        |               |             |
| LOCUS  | W60228   | 447 bp mRNA            | EST           | 15-OCT-1996 |
| DEFINITION   | Z328909_s1_Soares_fetal_heart_NbHH19W  | Homo sapiens           | CDNA clone    |             |
| IMAGE  | :42016 3'  | , mRNA sequence.       |               |             |
| ACCESSION  | W60228   |                        |               |             |
| NID  | 91366989   |                        |               |             |
| VERSION  | W60228.1   | GI:1366989             |               |             |
| KEYWORDS   | EST,   |                        |               |             |
| SOURCE   | human.   |                        |               |             |
| ORGANISM   | Homo sapiens   |                        |               |             |
| Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.   |  |                        |               |             |
| REFERENCE  | 1 (bases 1 to 447)   |                        |               |             |
| AUTHORS  | Hillier,L., Clark,N., Dubuge,T., Elliston,K., Hawkins,M., Holman,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J., Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston,R., Williamson,A., Wohldmann,P., Wilson,R.  |                        |               |             |
| TITLE  | The WashU-Merck EST Project  |                        |               |             |
| JOURNAL  | Unpublished (1995)   |                        |               |             |
| COMMENT  | On Apr 14, 1993 this sequence replaced gi:716583.  |                        |               |             |
| FEATURES   |  |                        |               |             |
| Source   | Contact: Wilson RK<br>Washington University School of Medicine<br>4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108<br>Tel: 314 286 1100<br>Fax: 314 286 1810<br>Email: est@watson.wustl.edu<br>This clone is available royalty-free through LILN; contact the IMAGE Consortium (info@image.liln.gov) for further information.<br>Insert Length: 857 Std Error: 0.00<br>Seq. Primer: mob:REGA+ET<br>High quality sequence stop: 414. |                        |               |             |
| Source   | Contact: Wilson RK<br>Washington University School of Medicine<br>4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108<br>Tel: 314 286 1100<br>Fax: 314 286 1810<br>Email: est@watson.wustl.edu<br>This clone is available royalty-free through LILN; contact the IMAGE Consortium (info@image.liln.gov) for further information.<br>Insert Length: 857 Std Error: 0.00<br>Seq. Primer: mob:REGA+ET<br>High quality sequence stop: 405. |                        |               |             |
| FEATURES   | Location/Qualifiers  |                        |               |             |
| Source   | 1. .447<br>/organism="Homo sapiens"<br>/db_xref="GDB:1.270.16"<br>/map="Q13; 754A01"; 12; 12q12-12q13.13"<br>/clone="IMAGE:345041"<br>/sex="unKnown"<br>/dev_stage="19 weeks"<br>/lab_host="DHL0B (ampicillin resistant)"  |                        |               |             |

/note="Organ: heart; Vector: pT7T3D (Pharmacia) with a modified polylinker; Site\_1: Not I; Site\_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dt) primer [5'-TGTTACCAATCTAAGGGGAGCGCCGGAAATTTTTTTTTTTTTTTTT T-3'], on equal amounts of mRNA from 2 13.5dpc and 2 14.5dpc embryos (total RNA provided by Minoru Ko, Wayne State Univ., from 2 ); double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and Eco RI adaptors (Pharmacia), digested with Not I and Eco RI sites of a modified pT7T3 vector (Pharmacia). Library went through one round of normalization, and was constructed by M. Fatima Bonaldo. This library was constructed from the same fetus as the fetal lung library, Soares fetal lung NbHL19W."

BASE COUNT 122 a 126 c 113 g 109 t

ORIGIN

Query Match 68.7%; Score 15.8; DB 26; Length 470;

Best Local Similarity 63.6%; Pred. No. 1.2e+02;

Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

Qy 2 crgtnqartggaytgcng 23

Db 435 CAGTTGGGGGACTTGGCCAG 414

RESULT 10 AA003148

LOCUS 324 bp mRNA EST

DEFINITION mg51601..r1 Soares mouse embryo NbMB13.5 14.5 Mus musculus cDNA

clone IMAGE:427344 5', mRNA sequence.

ACCESSION AA003148

VERSION 91446605

NID 1 AA003148..1 GI:1446605

KEYWORDS EST

SOURCE house mouse

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Bacteria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 324)

AUTHORS Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,,

Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,

Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,

Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and

Waterson,N.

TITLE The WashU-HMMI Mouse EST Project

JOURNAL Unpublished (1996)

COMMENT On Nov 4, 1993 this sequence version replaced.

Contact: Marra M/Mouse EST Project

WashU-HMMI Mouse EST Project

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: mouseest@watsen.wustl.edu

This clone is available royalty-free through LInL ; contact the

IMAGE Consortium (info@image.llnl.gov) for further information.

MGI: 261896 Seq primer: ETPrimer

High quality sequence stop: 287.

Location/Qualifiers

1. .324

High quality sequence stop: 287.

Location/Qualifiers

1. .324

High quality sequence stop: 287.

Location/Qualifiers

was primed with a Not I - oligo(dt) primer [5', TGTACCATCTAAGGGGAGCGCCGGAAATTTTTTTTTTTTTTTTT T-3'], on equal amounts of mRNA from 2 13.5dpc and 2 14.5dpc embryos (total RNA provided by Minoru Ko, Wayne State Univ., from 2 ); double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and Eco RI sites of a modified pT7T3 vector (Pharmacia). Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo. " M.Fatima Bonaldo, " was constructed by Bento Soares and normalization, and was constructed by Bento Soares and M. Fatima Bonaldo. This library was constructed from the same fetus as the fetal lung library, Soares fetal lung NbHL19W."

BASE COUNT 79 a 90 c 94 g 61 t

ORIGIN

Query Match 68.7%; Score 15.8; DB 27; Length 324;

Best Local Similarity 63.6%; Pred. No. 1.2e-02;

Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

Qy 2 cngtgartggaytgcng 23

Db 109 CAGTGAGGGGAACTTGGCCAG 130

RESULT 11 AA010775

LOCUS AA010775 432 bp mRNA EST

DEFINITION ze2zf04..r1 Soares\_fetal\_heart\_NbHL19W Homo sapiens cDNA clone IMAGE:359743 5', mRNA sequence.

ACCESSION AA010775

NID 91471802

VERSION AA010775..1 GI:1471802

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 432)

AUTHORS Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman,M., Hultman.M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J., Raffkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston,R., Williamson,A., Wohldmann,P., and Wilson,R.

TITLE The WashU-Merck EST Project

JOURNAL Unpublished (1995)

COMMENT On Apr 14, 1993 this sequence version replaced gi:785787.

Contact: Wilson RK Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: estwatson.wustl.edu This clone is available royalty-free through LInL ; contact the IMAGE Consortium (info@image.llnl.gov) for further information. Insert Length: 697 Std Error: 0.00 Seq primer: mob\_REGA+ET

FEATURES High quality sequence stop: 346.

Location/Qualifiers

1. .432

Source /organism="Homo sapiens"

/db\_xref="SDB:1276287"

/db\_xref="taxon:9606"

/map="4\_q35-qter"

/clone="IMAGE:359743"

/clone\_lib="Soares\_fetal\_heart\_NbHL19W"

/sex="unKnown"

/dev\_stage="19 weeks"

/lab\_host="DH10B (ampicillin resistant)"

/note="Organ: heart; Vector: pT7T3D (Pharmacia) with a modified polylinker; Site\_1: Not I; Site\_2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dt) primer [5', TGTACCATCTAAGGGGAGCGCCGGAAATTTTTTTTTTTTTTT T-3'], TGTACCATCTAAGGGGAGCGCCGGAAATTTTTTTTTTTTTTT T-3', double-stranded cDNA was size selected, ligated to Eco RI

adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT7T3 vector (Pharmacia). Library went through one round of normalization to a Cot = 5. Library constructed by M. Fatima Bonaldo. This library was constructed from the same fetus as the fetal lung library, Soares fetal lung NBHL19N."

BASE COUNT 108 a 113 c 105 g 106 t

## ORIGIN

Query Match 68.7%; Score 15.8; DB 27; Length 497;  
Best Local Similarity 63.6%; Pred. No. 1.2e+02; Indels 0; Gaps 0;  
Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

Qy 2 cngtngartgaaatytcnmg 23  
| || | : | || : | : || : |  
Db 215 CAGTGAGGGAACTTGTCCAG 236

RESULT 13  
AA036824

LOCUS AA036822 497 bp mRNA EST 26-AUG-1996  
DEFINITION IMAGE:471921 5', mRNA sequence.  
ACCESSION AA036822  
VERSION 91509860  
NID 214  
SOURCE AA036822.1 GI:1509860  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
Butheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 497)  
AUTHORS Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M.,  
Holman,M., Hultman,M., Kucaba,T., Le,M., Lenion,G., Marra,M.,  
Parsons,J., Raskin,L., Rohlfing,T., Soares,M., Tan,F.,  
Treviskis,E., Waterston,R., Williamson,A., Wohldmann,P. and  
Wilson,R.

TITLE The WashU-Merck EST Project  
JOURNAL Unpublished (1995)  
COMMENT On Apr 14, 1993 this sequence version replaced gi:785518.

Contact: Wilson RK  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: estwaton.wustl.edu  
Seq. primer: -28M13 rev2 from Amersham  
High quality sequence stop: 425.

FEATURES source  
1. 497

/organism="Homo sapiens"  
/db\_xref="SDB:3757923"  
/db\_xref="taxon:9606"  
/clone="IMAGE:471921"  
/clone\_id="Soares\_pregnant\_uterus\_NbHPtU"  
/sex="female"  
/dev\_stage="adult"  
/lab\_host="DH10B"

/note="Organ: uterus; Vector: PT7T3-PacI Site\_1: Not I - Site\_2: Eco RI; 1st strand cDNA was printed with a Not I - oligo(dt) primer [5'-TCATGGAAGAATTCCGGCGGCCATTTTTTTTTTTTTTTTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT7T3 vector. Library constructed by M. Fatima Bonaldo."

BASE COUNT 95 a 97 c 97 g 71 t

ORIGIN Query Match 68.7%; Score 15.8; DB 27; Length 360;  
Best Local Similarity 63.6%; Pred. No. 1.2e+02;

Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;  
Qy 2 cndtngartggatyytcnmg 23  
| || | : | || : | : || : |  
Db 172 CAGTGAGGGAACTTGTCCAG 193

RESULT 13  
AA038824

LOCUS AA038824 360 bp mRNA EST 28-AUG-1996  
DEFINITION m195907.r1 Soares mouse P3NMF19.5 Mus musculus cDNA clone  
IMAGE:474356 5', similar to WP:C28H8.12 CE01823 ; mRNA sequence.  
ACCESSION AA038824  
NID 91514249  
VERSION AA038824.1 GI:1514249  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus.  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
Butheria; Rodentia; Sciurognathi; Murinae; Mus.  
REFERENCE 1 (bases 1 to 360)  
AUTHORS Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,  
Geisler,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,  
Scheibenbogen,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,  
Theising,B., Wyllie,T., Lennon,G., Soares,B., Wilson,R. and  
Waterston,R.  
TITLE The WashU-HMM Mouse EST Project  
JOURNAL Unpublished (1996)  
COMMENT On Sep 21, 1992 this sequence version replaced gi:276467.

Contact: Marra M/Mouse EST Project  
WashU-HMM Mouse EST Project  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: mouseest@watson.wustl.edu

This clone is available royalty-free through LLNL ; contact the  
IMAGE Consortium (info@image.llnl.gov) for further information.  
Seq. primer: -28M13 rev2 from Amersham  
High quality sequence stop: 425.

FEATURES source  
1. 360  
/organism="Mus musculus"  
/clone\_id="Soares mouse P3NMF19.5"  
/map="8"  
/clone="IMAGE:474396"  
/db\_xref="taxon:10090"  
/note="vector: PT7T3D (Pharmacia) with a modified  
polylinker; Site\_1: Not I; Site\_2: Eco RI; 1st strand cDNA  
was primed with a Not I - oligo(dt) primer [5'-  
TGTTACAATCTGAGTGGAGGCCGGATTTTTTTTTTTTTTTTTT 3'],  
double-stranded cDNA was size selected, ligated to Eco RI  
adapters (Pharmacia), digested with Not I and cloned into  
the Not I and Eco RI sites of a modified pT7T3 vector  
(Pharmacia). Library went through one round of  
normalization to a Cot = 5. Library constructed by Bento  
Soares and M. Patima Bonaldo. RNA was kindly provided by  
Dr. Minoru Ko (Wayne State University)."

BASE COUNT 95 a 97 c 97 g 71 t

ORIGIN Query Match 68.7%; Score 15.8; DB 27; Length 360;  
Best Local Similarity 63.6%; Pred. No. 1.2e+02;

BASE COUNT 123 a 125 c 131 g 114 t 4 others

Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0; RESULT 15.

QY 2 cngtngartgaaattygcning 23 LOCUS WB2322 430 bp mRNA EST 12-SEP-1996  
Db 299 CAGTTGAGGGAACTTGCAG 320 DEFINITION mf04911.r1 Soares mouse p3NMF19.5 Mus musculus cDNA clone IMAGE 404132 5' similar to wp:C28H8.12 CE01823 ; mRNA sequence.

RESULT 14 ACCESSION W82322  
LOCUS AA057660 603 bp mRNA EST 19-MAY-1997  
DEFINITION 2193d09.s1 Stratagene corneal stroma (#937222) Homo sapiens cDNA clone IMAGE 512177 3' similar to TR:G1255188 G1255188 DYNAMITIN. ;  
NID 91540001  
VERSION W82322.1 GI:1540001  
SOURCE EST.  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
KEYWORDS EST.  
REFERENCE 1 (bases 1 to 430).  
AUTHORS Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T., Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Theising, B., Wyllie, T., Lennon, G., Soares, B., Wilson, R. and Waterston, R.  
TITLE The WashU-HMMI Mouse EST Project  
JOURNAL Unpublished  
COMMENT On Sep 12, 1996 this sequence version replaced gi:1393548.  
Contact: Marra M/Mouse EST Project  
WashU-HMMI Mouse EST Project  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: mouseest@watson.wustl.edu  
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.  
MGI:241900  
Seq primer: -28M13 rev2 from Amersham  
High quality sequence stop: 393.  
Location/Qualifiers 1..430

FEATURES source  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="IMAGE:404132"  
/clone\_lib="Soares mouse p3NMF19.5"  
/dev\_stage="19.5 dpc total fetus"  
/lab\_host="DH10B (ampicillin resistant)"  
/note="vector: pPT3D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was printed with a Not I - oligo(dT) primer [5',/map="CTGACATCTGAAGGGGGCGCATTTTTTTTTTTTTTTTTT 3']"  
/map="12"  
/db\_xref="GDB:3844653"  
/db\_xref="taxon:9606"  
/clone="IMAGE:512177"  
/clone\_lib="Stratagene corneal stroma (#937222)"  
/dev\_stage="76 years"  
/lab\_host="SOLR cells (kanamycin resistant)"  
/note="Organ: cornea; Vector: pBluescript SK+; Site\_1: EcoRI; Site\_2: XbaI; Cloned unidirectionally. Primer: Oligo dT. Corneal fibroblasts grown from explants, 76 years. Average insert size: 1.5 kb; Uni-ZAP XR Vector; ~5' adaptor sequence: 5' GAATTCGGACGAG 3' -3' adaptor sequence: 5' CTCGAGTTTTTTTTTTTTTTT 3' " 150 a 150 c 153 g 148 t 2 others  
BASE COUNT 113 a 112 c 118 g 87 t  
ORIGIN

Query Match 2 cngtngartgaaattygcning 23  
Best Local Similarity 68.7%; Score 15.8; DB 27; Length 430;  
Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;  
ORIGIN

QY 2 cngtngartgaaattygcning 23  
Best Local Similarity 63.6%; Score 15.8; DB 27; Length 603;  
Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;  
Db 371 CAGTTGGGGAACTTGCCAG 392  
Query Match 2 cngtngartgaaattygcning 23  
Best Local Similarity 63.6%; Score 15.8; DB 27; Length 430;  
Matches 14; Conservative 4; Mismatches 4; Indels 0; Gaps 0;  
Db 439 CAGTTGGGGAACTTGCCAG 418  
Search completed: October 1, 1999, 15:03:38  
Job time: 4382 sec



